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Amendments to the Claims:

This listing of claims will replace all prior versions, and listings, of claims in the application.

**Listing of Claims:** 

1.- 8. (canceled)

9. (currently amended) A isolated double protein mutant RecA protein comprising a

deletion truncation of at least 13 - 25 amino acids acid residues from the carboxyl terminus and

the an amino acid change from a glutamate to a basic amino acid at position 38.

10. (currently amended) The protein of Claim 9 wherein the truncation is 17 amino

acids acid residues are deleted truncated from the carboxyl terminus.

11. (currently amended) The protein of Claim 9 wherein the basic amino acid is to

lysine.

12. (currently amended) The protein of Claim 9 wherein the basic amino acid is to

arginine.

13. (currently amended) The protein of Claim 9 wherein 17 amino acid acids

residues are deleted truncated from the carboxyl terminus and the glutamate is changed to lysine,

as set forth in SEQ ID NO. 3.

14. (withdrawn) A polynucleotide sequence, as set forth in SEQ ID NO. 4, encoding

the protein of Claim 13.

15. (original) The protein of Claim 13 comprising an enhanced capacity to displace a

DNA binding protein as compared to wild-type RecA.

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16. (currently amended) The protein of Claim 15 wherein the <del>DNA binding</del> protein is a single stranded DNA binding protein, SSB.

- 17. (original) The protein of Claim 13 comprising an increased steady-state DNA binding capacity during a DNA strand exchange reaction as compared to wild-type RecA.
  - 18. (original) The protein of Claim 17 wherein the DNA is single-stranded.
  - 19. (original) The protein of Claim 17 wherein the DNA is double-stranded.
- 20. (original) The protein of Claim 19 wherein the double-stranded DNA is linear or circular.
- 21. (original) The protein of Claim 17 wherein the DNA strand exchange reaction is pH dependent.
- 22. (currently amended) The protein of Claim 21 wherein the DNA strand exchange reaction induces complete product formation at between a pH between of 8.0 to 9.0 7.5—9.5.
- 23. (currently amended) The protein of Claim 21 wherein the DNA strand exchange reaction induces complete product formation at a pH of 8.5 (± 1.0).
- 24. (original) The protein of Claim 17 wherein the DNA strand exchange reaction is Mg2+ concentration dependent.
- 25. (original) The protein of Claim 24 wherein the Mg2+ concentration is between 4mM 8mM.
  - 26. (original) The protein of Claim 24 wherein the Mg2+ concentration is 5mM.
- 27. (original) The protein of Claim 13 wherein the protein promotes an extended reaction, wherein the extended reaction is at least a three-strand exchange reaction.

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28. (canceled)

29. (withdrawn) A method of catalyzing in vitro homologous DNA pairing and DNA strand exchange reactions comprising providing a sufficient amount of the protein of Claim 1.

30. (withdrawn) A method of catalyzing in vitro homologous DNA pairing and DNA strand exchange reactions comprising providing a sufficient amount of the protein of Claim 9.

31. (withdrawn) A method of increasing recombination efficiency of homologous DNA pairing and DNA strand exchange reactions in a cell comprising supplying to the cell a sufficient amount of the protein of Claim 1.

32. (withdrawn) A method of increasing recombination efficiency of homologous DNA pairing and DNA strand exchange reactions in a cell comprising supplying to the cell a sufficient amount of the protein of Claim 9.

33.-34. (canceled)

35. (original) A kit comprising the protein of Claim 9.

36. (original) A kit comprising the protein of Claim 13.

37. (new) An isolated double mutant RecA protein comprising a truncation of up to 17 amino acid residues from the carboxyl terminus and an amino acid change from a glutamate to a basic amino acid at position 38.

38. (new) A kit comprising the protein of Claim 37.

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- 39. (new) An isolated double mutant RecA protein comprising a truncation of 17 amino acid residues from the carboxyl terminus and an amino acid change from a glutamate to a basic amino acid at position 38, wherein the basic amino acid is lysine, as set forth in SEQ ID NO: 3.
  - 40. (new) A kit comprising the protein of Claim 39.
- 41. (new) An isolated double mutant RecA protein comprising a truncation of at least 13 25 amino acid residues from the carboxyl terminus and an amino acid change from a glutamate to a basic amino acid at position 38, wherein the basic amino acid is lysine.
  - 42. (new) An kit comprising the protein of Claim 41.
- 43. (new) An isolated double mutant RecA protein comprising a truncation of at least 13-20 amino acid residues from the carboxyl terminus and an amino acid change from a glutamate to a basic amino acid at position 38.
- 44. (new) An isolated double mutant RecA protein comprising a truncation of at least 13 17 amino acid residues from the carboxyl terminus and an amino acid change from a glutamate to a basic amino acid at position 38.

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